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## Ny04 Full Avec Keygen Software REPACK

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Dependent on the knockdown  
efficiency. For a successful  
knockdown, plasmid amounts  
>5 µg/well should be used.  
Reference Moffat, M., Khoo, C.  
S. H., and Ghosh, R. (2002).  
Morpholino antisense  
oligonucleotides are powerful

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tools for gene function studies in fish. *Embo J.* 21:1530-1535.

Knockdown affects the developmental stages of the embryos at late stages.

Embryos injected with morpholino antisense oligo are fertile. Embryo fertility may be increased if the morpholino antisense oligo is co-injected with fertilized eggs. Females injected with the morpholino antisense oligo produce viable offspring. S2 cells were

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cultured as described in Methods. Cells were collected in presence of 0.5% sodium dodecyl sulfate (SDS). Whole-cell lysate was extracted in Laemmli sample buffer and subjected to polyacrylamide gel electrophoresis followed by Western blot analysis.

Endogenous laminin  $\gamma 1$  was specifically detected with anti-laminin  $\gamma 1$ . Inhibition of laminin  $\gamma 1$  chain splicing was accomplished by the use of a

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morpholino oligo complementary to a sequence in intron 2 (table 2) which is detected by the MabA13 antibody. The morpholino oligo was designed to inhibit splicing at the 5'-splice site (first arrow) and the branchpoint of intron 2 (second arrow). The blocking of these splicing events results in the retention of the mRNA intermediate and an increase in the level of laminin  $\gamma$ 1 mRNA. Knockdown of laminin

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$\gamma 1$  results in a reduction of the expression level of the docked form of laminin  $\gamma 1$  and a concomitant increase in the expression level of the undocked form of laminin  $\gamma 1$  which is detected by the MabA13 antibody. This MabA13 antibody can be used for subsequent experiments on laminin  $\gamma 1$  expression regulation. For example  
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